

VESSEL CUTTING DEVICES

Cross Reference to Related Applications

This is a continuation of application  
No. 09/850,021, filed May 7, 2001, which is a  
5 continuation of application No. 09/014,759 (now U.S.  
patent 6,416,527), filed January 28, 1998, both of  
which are hereby incorporated by reference herein in  
their entireties.

Background of the Invention

10 This invention relates to vessel cutting  
devices for use in the repair, replacement or  
supplement of a medical patient's natural body organ  
structures or tissues. More particularly, this  
invention relates to vessel cutting devices for use in  
15 vascular anastomosis (the surgical connection of  
vessels).

An example of the possible uses of the  
invention is a minimally invasive cardiac bypass  
procedure. This and other examples are considered in  
20 detail in Goldsteen et al. U.S. patent 5,976,178, which  
is hereby incorporated by reference herein in its  
entirety.

Vascular anastomosis is a delicate and time-consuming procedure in which fast and accurate vessel cutting plays a particularly important role.

In view of the foregoing, it would be  
5 desirable to provide a catheter-based system for accessing specific body cavities percutaneously, thereby minimizing patient trauma.

It would also be desirable to provide fast and accurate vessel cutting devices.

10 Summary of the Invention

It is an object of the present invention to provide a catheter-based system for accessing specific body cavities percutaneously, thereby minimizing patient trauma. It is also an object to provide fast  
15 and accurate vessel cutting devices.

These and other objects are accomplished by providing a method and apparatus for creating an aperture at an access site in a patient's existing tubular body organ structure by passing a delivery  
20 sheath axially along the interior of a portion of the existing tubular body organ structure to place a distal end of the delivery sheath near the access site, passing a centering wire axially along the interior of the delivery sheath, piercing through from inside to  
25 outside of the patient's existing tubular body organ structure at the access site by causing an end portion of the centering wire to emerge from the distal end of the delivery sheath, passing a cutting catheter substantially coaxially over the centering wire and  
30 axially along the interior of the delivery sheath, forming the aperture by advancing a distal end of the cutting catheter through from inside to outside of the

patient's existing tubular body organ structure at the  
access site and advancing the distal end of the  
delivery sheath through from inside to outside of the  
patient's existing tubular body organ structure at the  
5 access site.

In one embodiment, the distal end of the  
cutting catheter is rotated to cut through the  
patient's existing tubular body organ structure at the  
access site. In another embodiment, a cutting catheter  
10 with a conical (preferably star-shaped) cutting edge is  
pushed through the patient's existing tubular body  
organ structure at the access site.

The present invention can also be used to  
create an aperture in the patient's existing tubular  
body organ structure by advancing a distal end of the  
cutting catheter through from outside to inside of the  
15 patient's existing tubular body organ structure at the  
access site.

In the most preferred embodiment, all or  
substantially all necessary apparatus is inserted into  
the patient via the patient's existing body organ  
vessel. In addition, all or substantially all  
apparatus functions are controlled by the physician (a  
term used herein to also include supporting  
25 technicians) from outside the patient's body.

#### Brief Description of the Drawings

The above and other objects and advantages of  
the invention will be apparent upon consideration of  
the following detailed description, taken in  
30 conjunction with the accompanying drawings, in which  
like reference characters refer to like parts  
throughout, and in which:

FIG. 1a is a simplified sectional view showing the distal end of a delivery sheath in the interior of a portion of the existing tubular body organ structure with a centering wire piercing through  
5 from inside to outside of the patient's existing tubular body organ structure at the access site;

FIG. 1b is a view similar to portions of FIG. 1a showing a centering wire piercing through from inside to outside of the patient's existing tubular  
10 body organ structure at the access site, wherein the end portion of the centering wire includes a selectively enlargeable structure;

FIG. 1c is another view similar to portions of FIG. 1a showing a centering wire piercing through  
15 from inside to outside of the patient's existing tubular body organ structure at the access site, wherein the end portion of the centering wire includes fasteners;

FIG. 2 is yet another view similar to FIG. 1a showing a cutting catheter positioned for cutting at  
20 the distal end of a delivery sheath at the access site;

FIG. 3 is still another view similar to FIG. 1a showing forming the aperture by advancing a distal end of the cutting catheter through from inside to  
25 outside of the patient's existing tubular body organ structure at the access site;

FIG. 4 is yet another view similar to FIG. 1a showing advancing the distal end of the delivery sheath through from inside to outside of the patient's  
30 existing tubular body organ structure at the access site;

FIG. 5 is a simplified elevational view, partly in section, showing the distal end of the

cutting catheter advancing through from outside to inside to create an aperture in the patient's existing tubular body organ structure;

FIG. 6 is a side view of the patient's  
5 existing tubular body organ structure of FIG. 5, showing the aperture created;

FIG. 7a is still another view similar to FIG. 1a showing the distal end of a delivery sheath in the interior of a portion of the existing tubular body  
10 organ structure with a centering wire piercing through from inside to outside of the patient's existing tubular body organ structure at the access site, wherein the cutting catheter includes a dilator;

FIG. 7b is yet another view similar to FIG. 1a forming the aperture by advancing a distal end of the cutting catheter through from inside to outside of the patient's existing tubular body organ structure at the access site, wherein the cutting catheter includes a dilator;

20 FIG. 7c is still another view similar to FIG. 1a showing advancing the delivery sheath through the aperture at the access site; and

FIG. 8 is yet another view similar to FIG. 1a showing a delivery sheath which includes distal and  
25 proximal selectively enlargeable structures.

#### Detailed Description of the Invention

As a preliminary step in creating an aperture at an access site 10 in a patient's existing tubular body organ structure 1, a delivery sheath 20 is passed  
30 axially along the interior of a portion of tubular body organ structure 1 to place a distal end of delivery sheath 20 near access site 10. When the distal end of

delivery sheath 20 is adjacent to access site 10, a centering wire 30 is passed axially along the interior of the sheath until the end portion of centering wire 30 emerges from the distal end of the sheath and pokes  
5 through from inside to outside of tubular body organ structure 1. Centering wire 30 provides a pilot track for cutting catheter 40 to follow. FIG. 1a shows the distal end of delivery sheath 20 in the interior of a portion of tubular body organ structure 1 with a  
10 centering wire 30 piercing through from inside to outside of the organ structure at access site 10.

The distal end of centering wire 30 is preferably deformable to facilitate deployment and removal, but resumes its operational (preferably  
15 hooked) shape once deployed. Centering wire 30 is kept relatively straight when it is inside sheath 20. But, when centering wire 30 is pushed axially out the distal end of sheath 20, it curves to one side, as shown in FIGS. 1a, 1b and 1c. FIGS. 1b and 1c show alternative  
20 structures for centering wire 30. In FIG. 1b, the end portion of centering wire 30 includes a selectively enlargeable structure (such as a balloon 50 which extends annularly around the exterior of the centering wire and projects radially outwardly from the centering  
25 wire in all radially outward directions when inflated). In FIG. 1c, the end portion of centering wire 30 includes struts 55 spaced circumferentially around centering wire 30 and which are resiliently biased to project from the centering wire after the end portion  
30 of the centering wire pierces through body organ structure 1 at access site 10. By providing a selectively enlargeable structure disposed on the exterior of the centering wire at a predetermined

distance proximally from the distal end of the centering wire and enlarging that structure after the centering wire has pierced organ structure 1, it is possible to prevent the portion of centering wire 30  
5 which is distal of the enlargeable structure from passing back into the organ structure. In addition to the retaining function, the enlargeable structure serves to seal the aperture and displace tissue from around the outside of organ structure 1 near access  
10 site 10, thereby creating a space. Such a space helps to prevent cutting head 45 from cutting other tissues after exiting organ structure 1 at access site 10.

After piercing through organ structure 1 at access site 10 with centering wire 30, cutting  
15 catheter 40 is passed substantially coaxially over the centering wire and axially along the interior of sheath 20. FIG. 2 shows cutting head 45 of cutting catheter 40 positioned for cutting at the distal end of delivery sheath 20 at access site 10.

20 Centering wire 30 holds cutting catheter 40 and delivery sheath 20 against organ structure 1 at access site 10, thereby preventing undue bleeding during and after the creation of the aperture that could occur if the cutting catheter and the delivery  
25 sheath were to move away from the access site. FIG. 3 shows how the aperture is formed by advancing the distal end of cutting catheter 40 (i.e., cutting head 45) through from inside to outside of organ structure 1 at access site 10 by rotating and/or pushing the distal  
30 end of the cutting catheter.

As shown in FIGS. 2, 3, and 4, the distal end of cutting catheter 40 has a circular cutting edge. Cutting catheter 40, which when advanced by rotation,

cuts through tissue and removes tissue plug 60. The preferred embodiment of cutting head 45 also includes a serrated cutting edge and an axially aligned recess for accepting tissue plug 60. By removing plug 60 of  
5 tissue (rather than merely displacing tissue, as in FIGS. 5 and 6), the elastic recoil of organ structure 1 at access site 10 is reduced, which may be a desirable condition for optimal graft attachment.

FIG. 4 shows advancing the distal end of  
10 delivery sheath 20 through from inside to outside of organ structure 1 at access site 10 and removing centering wire 30 and cutting catheter 40 along with tissue plug 60 contained within cutting head 45.

As shown in FIG. 5, non-rotating cutting  
15 catheter 40 can be used to create specific geometric aperture shapes (e.g., oblong aperture 70 for coronary anastomosis). FIG. 5 also shows the use of the present invention in creating an aperture in organ structure 1 by advancing a distal end of cutting catheter 40  
20 through from outside to inside of the organ structure at access site 10. Centering wire 30 is tracked through cutting catheter 40 and is shown piercing organ structure 1 at access site 10. Following such an outside-to-inside aperture, delivery sheath 20 can be  
25 passed axially along the interior of a portion of organ structure 1 to place a distal end of delivery sheath 20 near second access site 10 where an inside-to-outside aperture can be created. (Note that organ structure 1 is shown smaller in scale relative to sheath 20 and  
30 cutting catheter 40.)

FIG. 6 is a side view of organ structure 1, showing aperture 70 created using non-rotating cutting catheter 40 of FIG. 5.



Cutting catheter 40 shown in FIG. 7a is a rotating catheter. Cutting head 45 could be a saw-tooth or a razor-edge type, for example. The distal end of delivery sheath 20 is shown in the interior of a portion of organ structure 1 with centering wire 30 piercing through from inside to outside of the organ structure at access site 10, wherein cutting catheter 40 includes dilator 80. Dilator 80 facilitates advancing sheath 20 through the aperture (as is shown by the succession of steps illustrated by FIGS. 7b and 7c).

The outer diameter of dilator 80 is close to the inner diameter of sheath 20 and is typically larger than the diameter of cutting head 45. As shown in FIG. 7b, as dilator 80 advances through the aperture at access site 10, the aperture is simultaneously sealed against bleeding.

FIG. 8 shows delivery sheath 20 which includes proximal and distal selectively enlargeable structures 90, 100. Preferably, both selectively enlargeable structures 90, 100 are balloons which extend annularly around the exterior of delivery sheath 20 and project radially outward when inflated. Although the embodiment shown in FIG. 8 includes both proximal and distal selectively enlargeable structures, either one or both may be included. When enlarged, proximal selectively enlargeable structure 90 prevents more than the portion of delivery sheath 20 which is distal of the enlargeable structure from passing out of the tubular structure via the aperture. Similarly, when enlarged, distal selectively enlargeable structure 100 prevents the portion of delivery sheath 20 which is

distal of the enlargeable structure from passing back in to the tubular structure via the aperture.

As an illustrative example of the application of the present invention, consider the following.

- 5 Delivery sheath 20 (preferably about 4.0 mm in diameter) including cutting catheter 40 is introduced into organ structure 1 percutaneously through the femoral artery near the thigh. Cutting catheter 40 includes cutting head 45 (preferably about 3.5 mm in  
10 diameter). Delivery sheath 20 is positioned at access site 10, here the ascending aorta. Centering wire 30 is tracked through cutting catheter 40 and is caused to pierce the aortic artery at access site 10. Cutting catheter 40 is then tracked over centering wire 30 by  
15 either pushing or rotating (or a combination of both pushing and rotating) and caused to advance through the aortic wall. An approximately 3.5 mm aperture is created with tissue plug 60 retained in cutting head 45 and removed along with the cutting catheter 40.
- 20 Delivery sheath 20 can now be advanced through the approximately 3.5 mm aperture created by the cutting catheter 40, causing organ structure 1 to stretch slightly (i.e., about 0.5 mm). This stretching is desirable because it provides a blood seal around  
25 delivery sheath 20 to prevent bleeding into the chest cavity. Delivery sheath 20 can now be used to introduce other catheters (including cameras, for example) from the femoral artery into the chest cavity for the purpose of diagnosis or intervention (e.g.,  
30 grafts or TMR laser surgery).

To minimize patient trauma, delivery sheath 20, cutting catheter 40 and centering wire 30

are all preferably coupled to and controlled by a controller located on the outside of the patient.

Various methods and apparatus for delivering and installing plugs in walls of organ structures, as  
5 well as methods and apparatus for promoting the closing and healing of apertures in walls of organ structures, are available (e.g., of the type shown in Goldsteen et al. U.S. patent 5,976,178; Goldsteen et al. published PCT patent application WO 98/47430; and  
10 Sullivan et al. published PCT patent application WO 98/55027, all of which are hereby incorporated by reference herein).

Thus, it is seen that a method and apparatus for creating an aperture at an access site in a  
15 patient's existing tubular body organ structure and making it possible to access specific body cavities percutaneously, thereby minimizing patient trauma, is provided. One skilled in the art will appreciate that the present invention can be practiced by other than  
20 the described embodiments, which are presented for purposes of illustration and not of limitation, and the present invention is limited only by the claims which follow.